## **Device Registration and Listing Module**

Form Number: FDA 3673(03/08)

OMB Number: 0910-0625

OMB Expiration Date: xx/xx/20xx

**OMB Burden Statement:** 

Public reporting burden for this collection of information on form FDA 3673 is estimated to be 0.50 hours per response for the purpose of firms annually registering their establishment and 0.25 hours per response for the purpose of firms annually listing their devices. These estimates are based on FDA's experience, data from the device registration and listing database, and our estimates of the time needed to complete other previously required forms.

Send comments regarding this burden estimate or another aspect of this collection of information, including suggestions for reducing this burden to:

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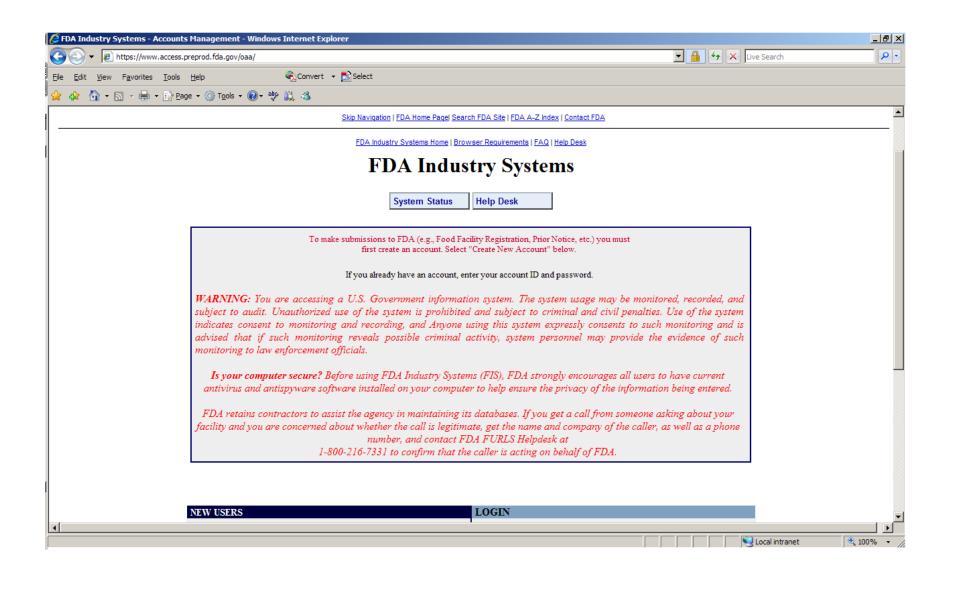
Food and Drug Administration,

8455 Colesville Rd.,

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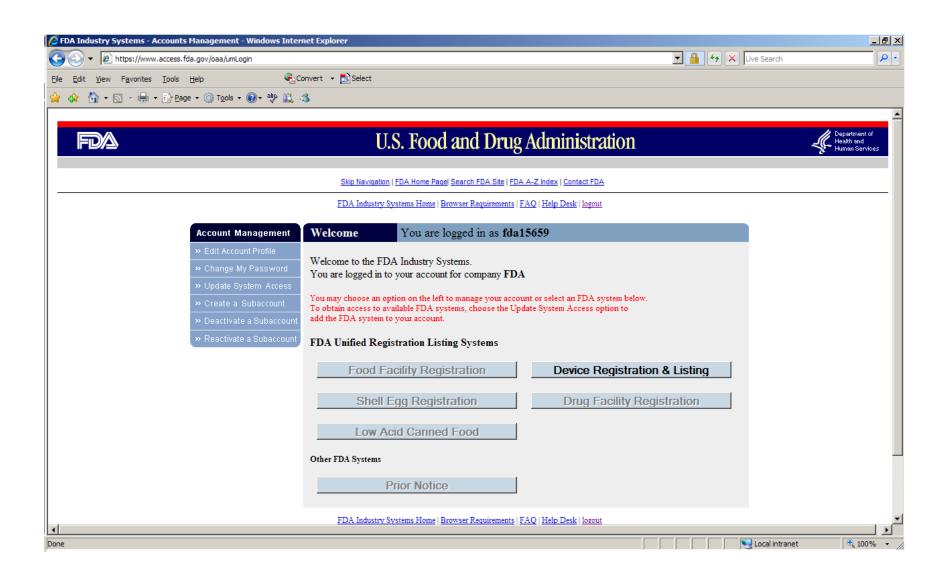


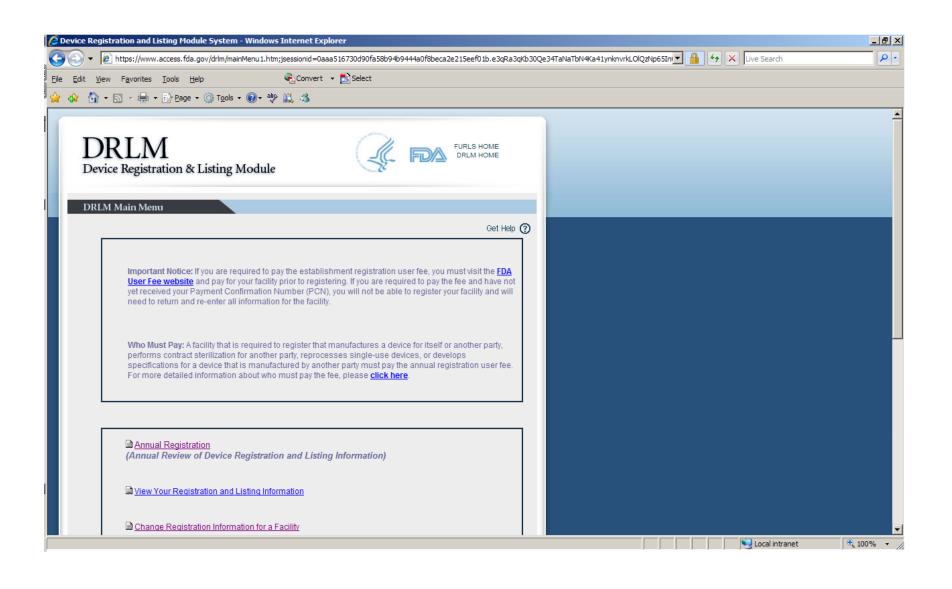
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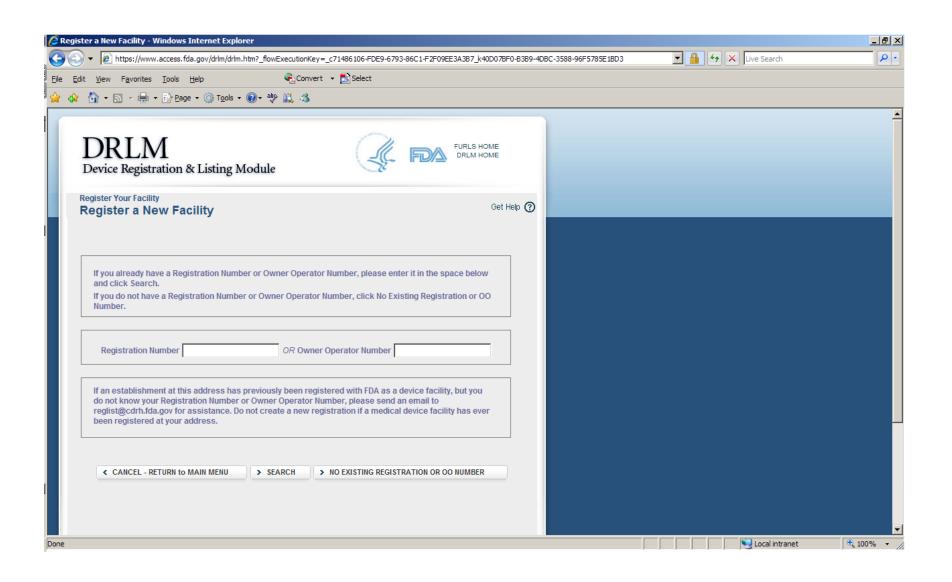
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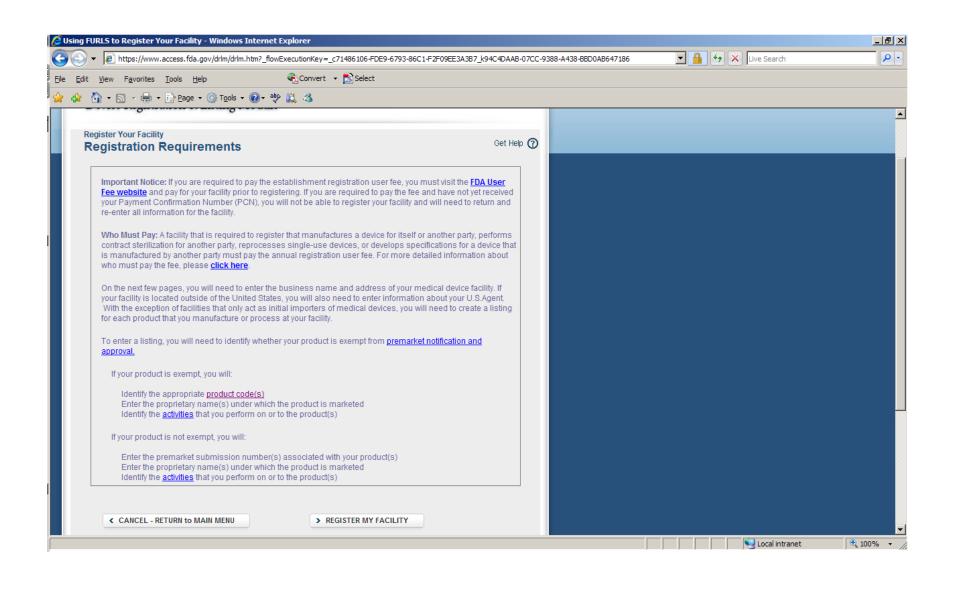


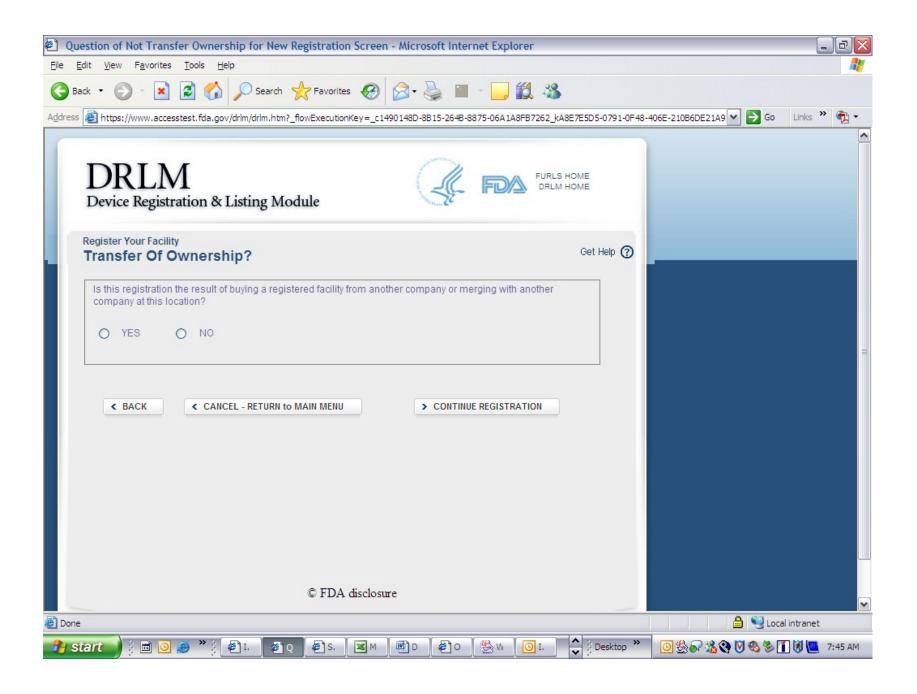


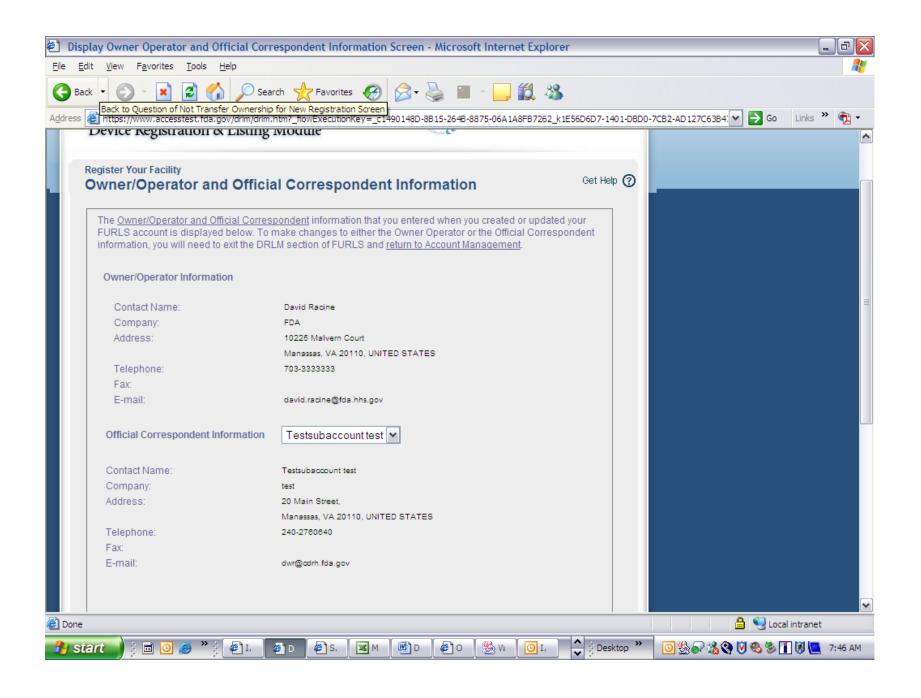
- a Cancel, Deactivate, or Reactivate a Facility Registration
- Change the Official Correspondent for a Facility
- Register a New Medical Device Facility
- Create Listings for Medical Devices
- Change, Cancel, or Reactivate Listings
- Transfer Ownership of Devices or Facilities

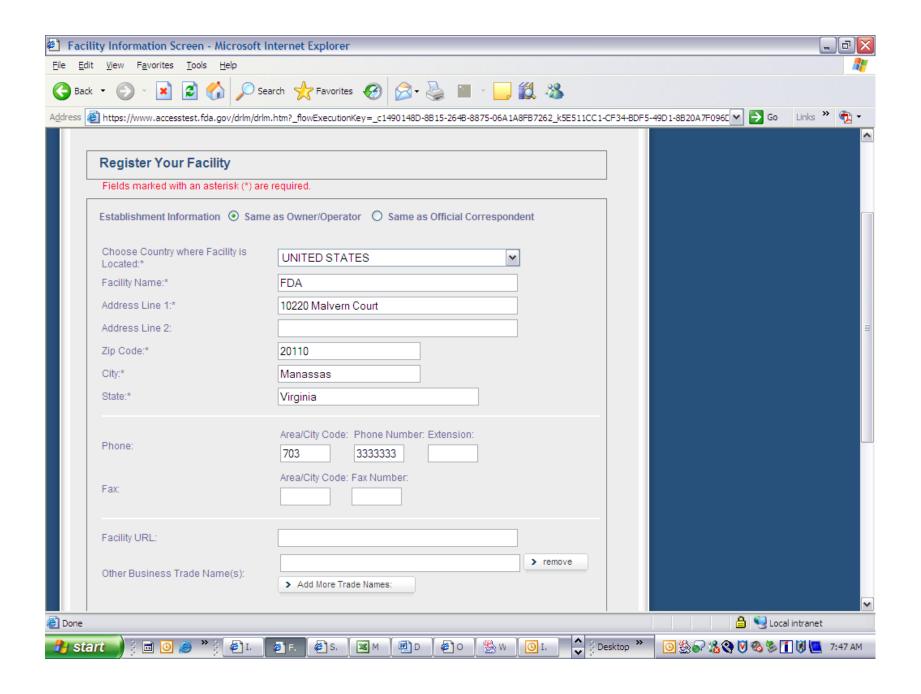
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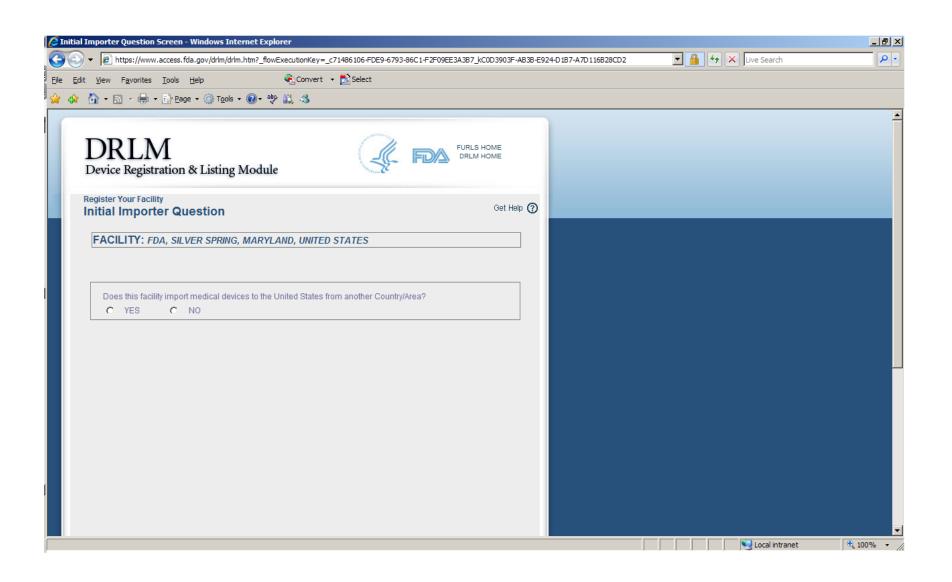


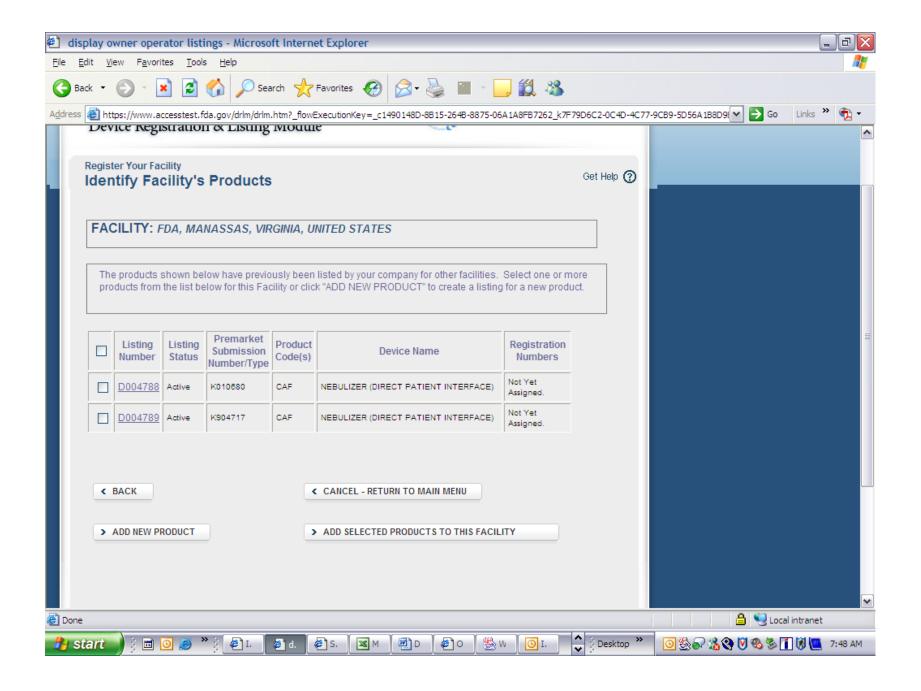


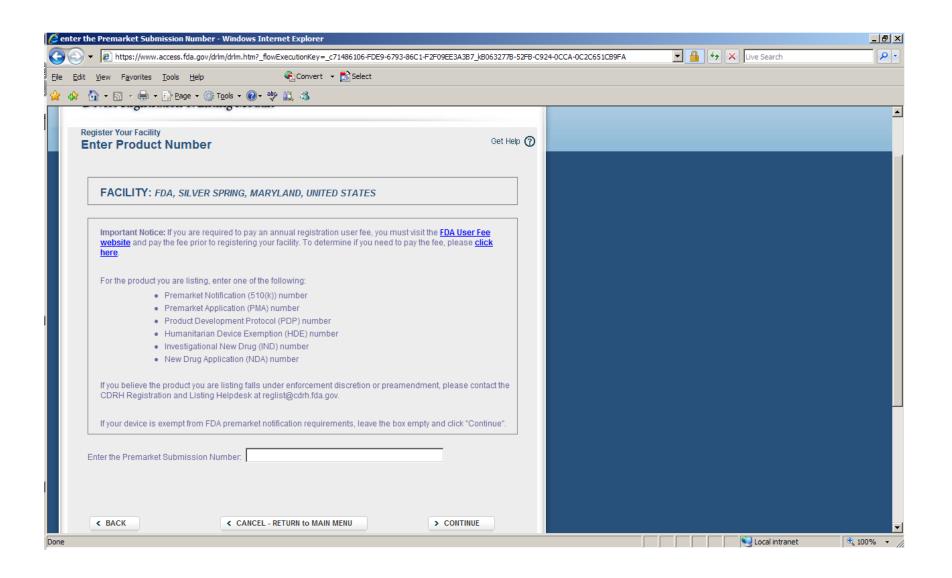


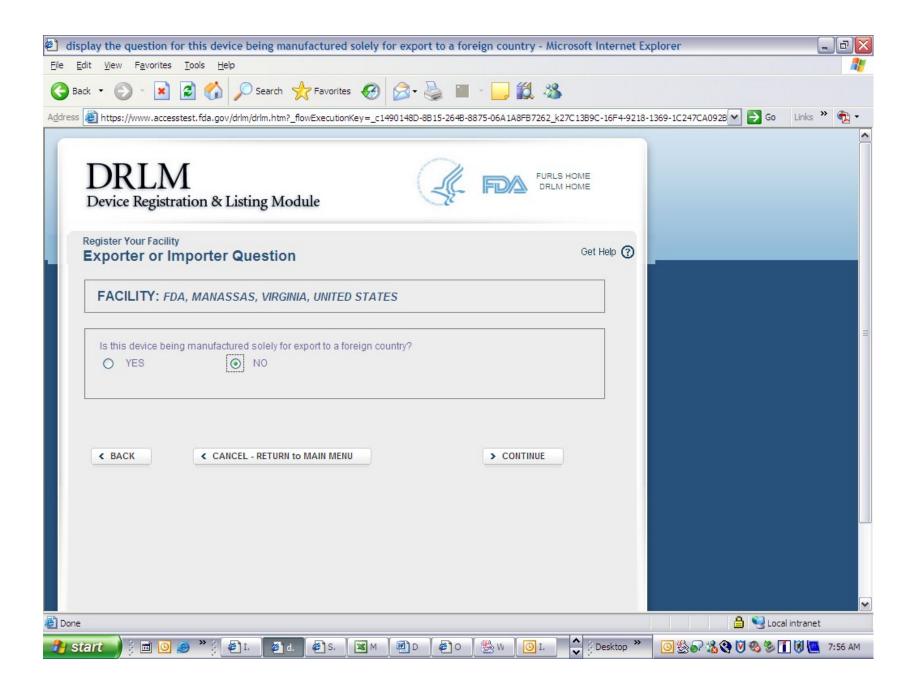


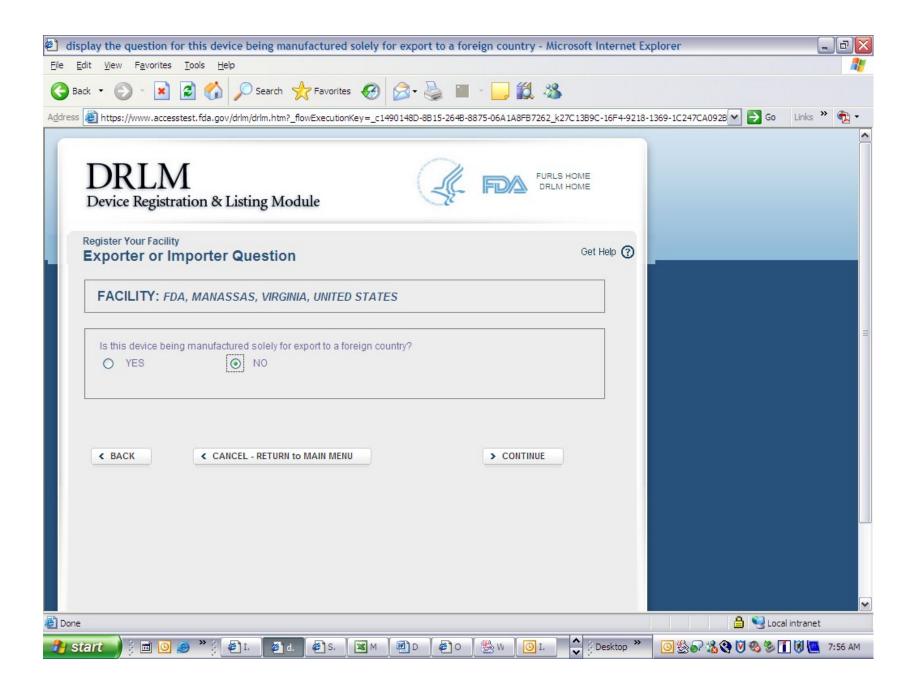


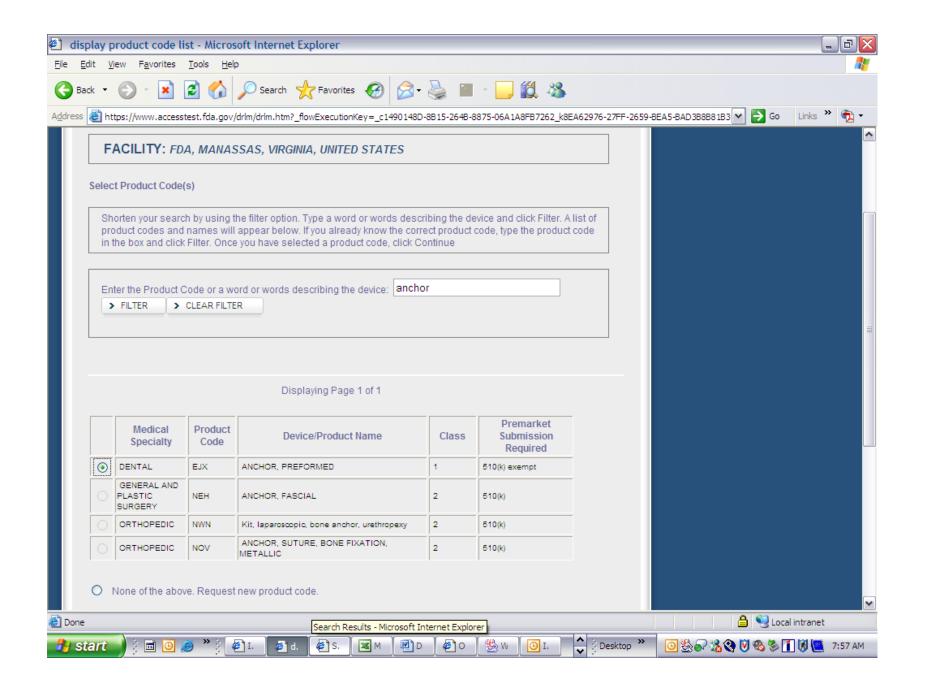


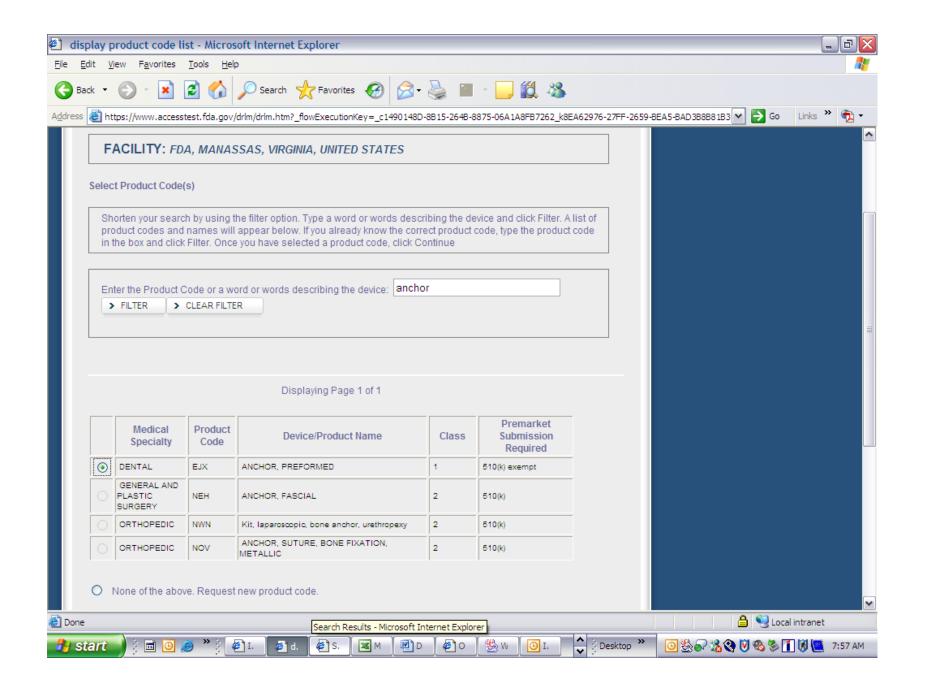


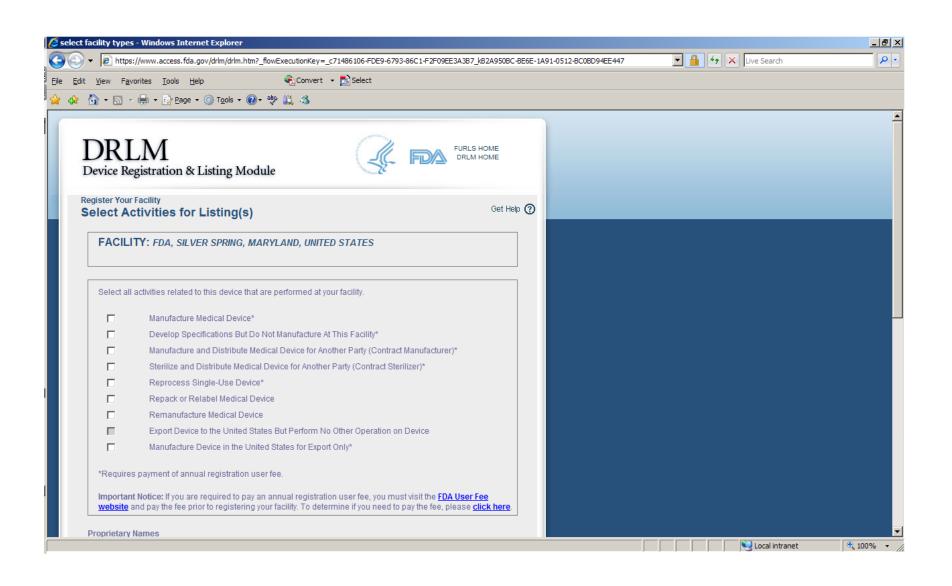




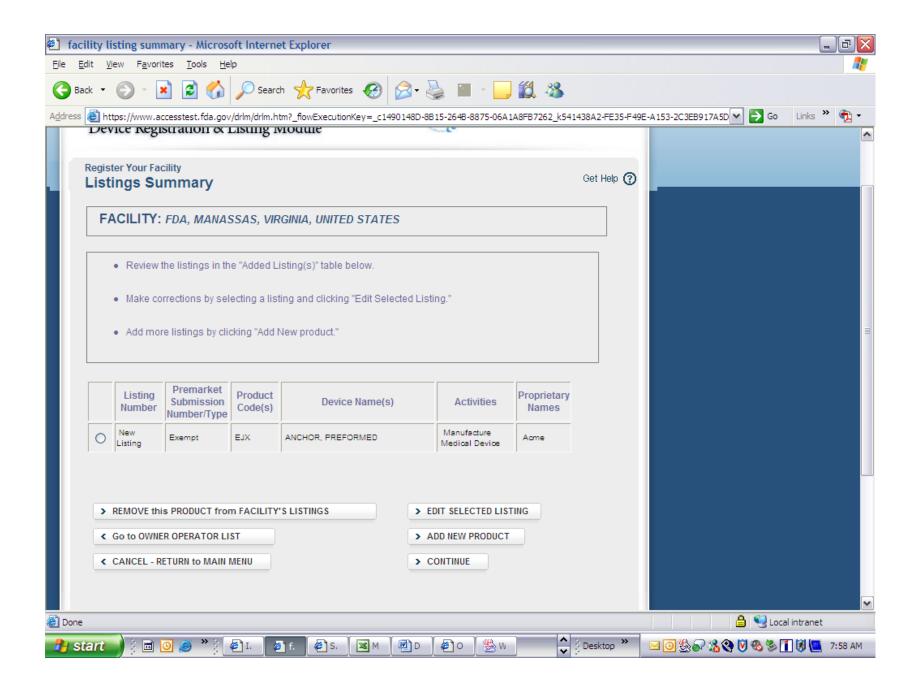


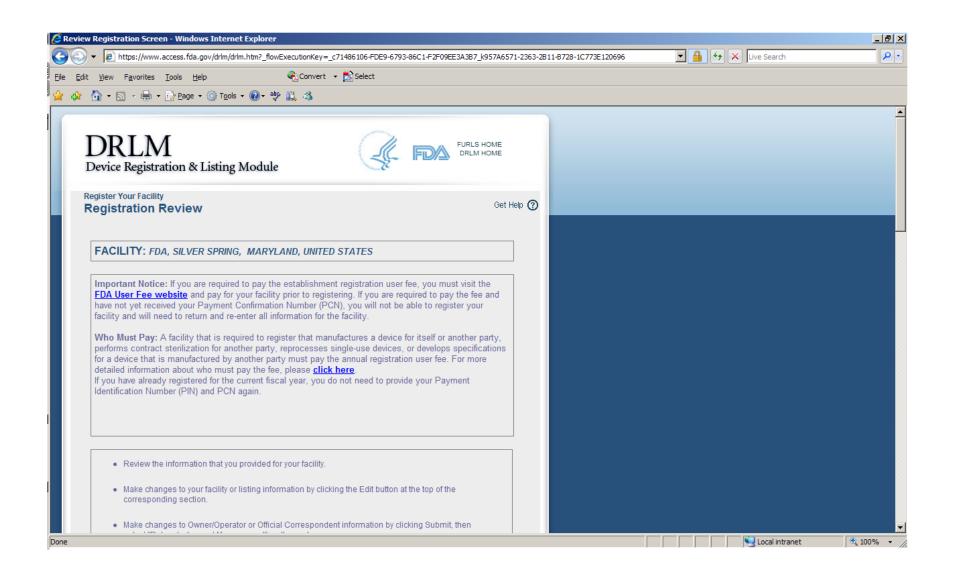


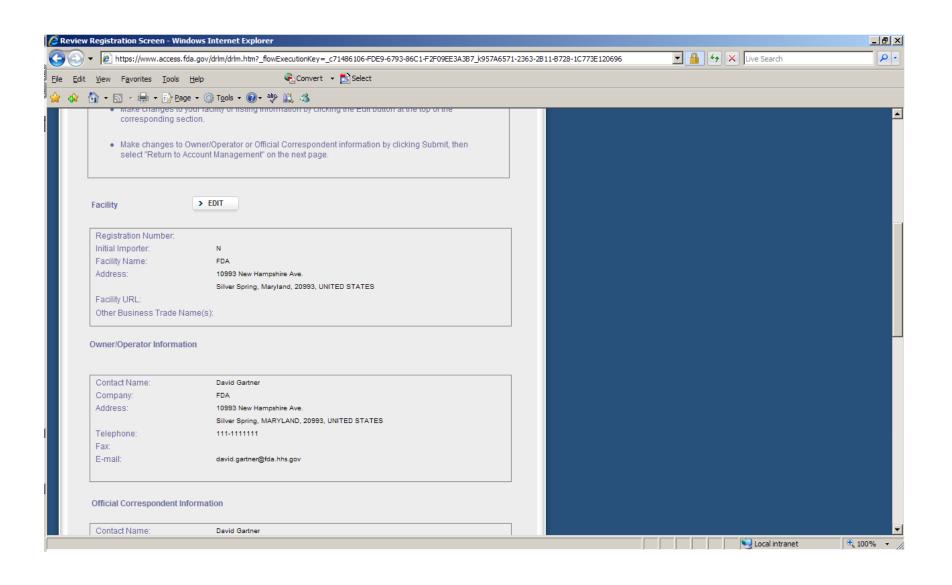


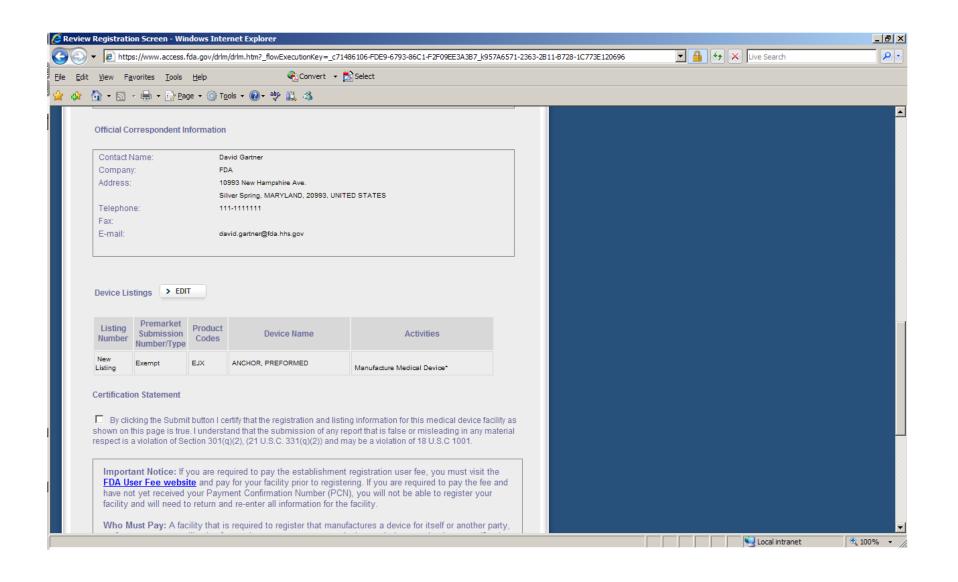


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Who Must Pay: A facility that is required to register that manufactures a device for itself or another party, performs contract sterilization for another party, reprocesses single-use devices, or develops specifications for a device that is manufactured by another party must pay the annual registration user fee. For more detailed information about who must pay the fee, please <u>click here</u>.

If you have already registered for the current fiscal year, you do not need to provide your Payment Identification Number (PIN) and PCN again.

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